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Applicant: **VENTRITEX, INC.**
701 E. Evelyn Avenue
Sunnyvale,
California 94025 (US)

Inventor: **Fain, Eric S.**
236 Marmona Drive
Menlo Park, CA 94025 (US)

Representative: **Witte, Alexander, Dr.-Ing. et al**
Witte, Weller, Gahlert & Otten
Patentanwälte
Rotebühlstrasse 121
D-70178 Stuttgart (DE)

Method and apparatus for delivering defibrillation shocks with improved effectiveness.

A method for defibrillating a patient's heart with an implantable defibrillator having defibrillation synchronized to the fibrillation waveform is disclosed. The defibrillator is used to detect fibrillation in a patient's heart, monitor at least one far-field fibrillation voltage across two spaced-apart electrodes while the high voltage capacitors are charging, then deliver a high energy shock during a period of ventricular fibrillation when the absolute values of the peak and trough voltages exceed a threshold based

on a desired percentage greater than a running average of the absolute values of the peaks and troughs of the fibrillation voltages monitored during and after capacitor charging. The shock is then delivered synchronous with the next significant cardiac complex. The defibrillator also includes a safety timer so that if the threshold is not reached within a specified safe time limit, the energy shock will be delivered without further delay.

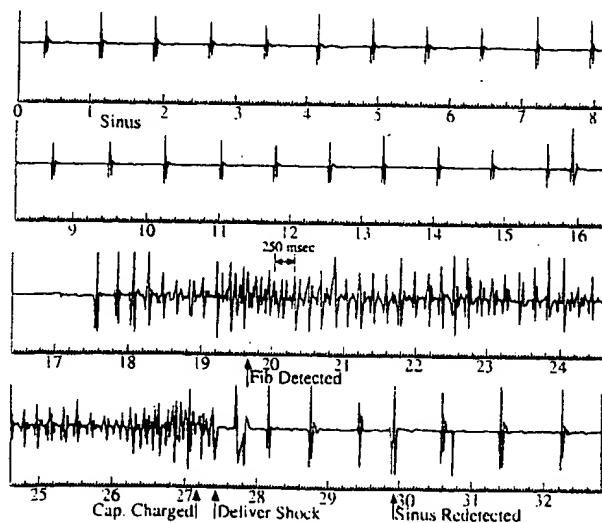


Figure 3

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FIELD OF THE INVENTION

The present invention relates generally to implantable cardiac stimulation devices, and more specifically to an implantable defibrillator that delivers defibrillation energy synchronously with the fibrillation waveform.

BACKGROUND OF THE INVENTION

Following detection of ventricular fibrillation, prior art implantable defibrillators typically deliver a defibrillation shock as soon as their high voltage (HV) capacitors are charged, and in more advanced systems, after ventricular fibrillation (VF) is reconfirmed. The timing of the shock is either not related to the status of the VF signal or is delivered synchronous with the next cardiac complex after reconfirmation and/or charge completion.

In "Genesis of Sigmoidal Dose-Response Curve During Defibrillation by Random Shock: A Theoretical Model Based on Experimental Evidence for a Vulnerable Window During Ventricular Fibrillation," in *PACE* 1990, 13:1326-1342, Hsia and Mahmud discuss a theoretical model and experimental results for using a surface electrocardiogram (ECG) to identify a window of VF during which the heart is more susceptible to defibrillation. In their experiments, they compared the VF waveform voltage at the time of shock delivery for successful and unsuccessful shocks. The shock strength was held constant, and was chosen to produce a defibrillation success rate of 50%. When using a recording from a lead II surface ECG, the absolute VF voltage (AVFV) was found to exhibit significantly larger values for successful defibrillation as compared to unsuccessful defibrillation. However, their experiments using intracardiac electrograms from two epicardial patch electrodes instead of a lead II ECG to track the state of VF showed no significant difference in AVFV between successful and unsuccessful defibrillation. They do not disclose how using the window of VF susceptible to defibrillation could be used in an implantable device.

In an abstract from an American Heart Association conference in Atlanta, November 1993, Hsia and Frerk et al. describe "On-line Electronic Identification of a Period of Vulnerability to Defibrillation using Real-Time Ventricular Fibrillation Waveform Analysis". If implemented as described, however, this system would not work in a conventional implantable device because the electrogram from a bipolar pair of endocardial pacing and sensing electrodes would not provide the information necessary to generate a useful AVFV, which is needed to determine the window of susceptibility to defibrillation.

SUMMARY OF THE INVENTION

The implantable defibrillator of the present invention reduces the defibrillation threshold (DFT) by delivering a defibrillation shock during a period of ventricular fibrillation (VF) when the heart is most susceptible to defibrillation. Variation in the state of excitability of cardiac tissue results in a cyclic period of increased susceptibility to defibrillation. The susceptible period occurs at a time when the number of excitable cells is low, i.e., a higher state of total depolarization. Far-field sensing is used to provide an electrogram that is monitored during VF, beginning with charging of the high voltage capacitors, to determine a relative measure of the absolute ventricular fibrillation voltage (AVFV), which is a crude marker of the number of excitable cells. Because an electrogram is sensitive to the location of the sensing electrodes, it is important to use unipolar, i.e. far-field, sensing to find the higher state of total depolarization in the heart. A bipolar signal, which is the usual signal used for sensing in implantable defibrillators, gives information regarding only the VF wavefronts passing through the localized area between the two electrodes, typically the RV pacing and RV sensing electrodes. The amplitude of the signal depends, in part, on the direction the wavefront is traveling; the signal will be largest when traveling parallel to the two electrodes, and smallest when perpendicular to them. By using electrodes that are at least 4 centimeters apart, a more global ventricular fibrillation signal is obtained. This signal is continually monitored, usually beginning with charging of the high voltage capacitors. In the preferred embodiment, a running average of the AVFV, for example the last 5 seconds, is utilized to determine whether the ECG signal is relatively large or small. When the capacitors are fully charged, the shock is delivered during a period of VF when the absolute values of the peak and trough voltages, calculated within a successive series of 250 msec windows exceed the calculated average monitored during the running window by a desired amount, for example 125%. The running average is calculated by looking at a predetermined number of the most recent 250 msec windows, typically 5 seconds or about 20 windows. The value of each window may be calculated by averaging the absolute value of all the data points in a window, averaging the absolute values of all the peaks and troughs in a window or looking at the area under the curve for the window. In the preferred embodiment only the most recent signal is considered since older data may not be relevant in determining when to deliver the defibrillation shock. The defibrillation shock is timed to coincide with a relative maximum absolute VF voltage (a peak or a trough).

In an alternative embodiment, a relative maximum amplitude of the absolute VF voltages from the windows is calculated and a shock is delivered upon the completion of charging of the high voltage capacitors and reconfirmation of VF when the measured AVFV is within a desired percentile of the relative maximum, for example 50 to 90%. As with the running average, the number of windows used to determine the relative maximum is limited to the most recent time period, for example 5 seconds. This ensures that only the most relevant portion of the VF signal is used to determine when to deliver the defibrillation shock.

More than two electrodes may be used in combination to generate the VF signal or signals. Two pairs of electrodes may be used, with or without one electrode in common, to observe fibrillation wavefronts from two directions.

The electrodes used for VF monitoring may be the same electrodes used to deliver the high energy defibrillation shock and provide pacing and sensing. This is advantageous since no extra leads are required, and the present invention can be used with existing lead systems.

A safety timer is included in the defibrillator so that if a relatively large absolute VF voltage is not obtained to trigger delivery of the shock within a certain time, say 5 seconds from the completion of charging of the high voltage capacitors and reconfirmation of VF, the shock will be delivered. The shock will still be delivered synchronous with the next cardiac complex. Note that the safety timer could alternatively be set to time from the detection of initiation of VF.

The present invention is directed toward the provision of a defibrillator that increases the likelihood for successful defibrillation at any particular shock strength by delivering the defibrillation pulse during a period corresponding to a relatively high state of depolarization of the heart cells as monitored by a unipolar electrogram during VF.

It is thus a primary object of this invention to provide an implantable cardiac defibrillator that delivers a defibrillation pulse at a time most likely to defibrillate the heart.

It is a further object of this invention to provide an implantable cardiac stimulation system that can be used with previously implanted defibrillation lead systems.

BRIEF DESCRIPTION OF THE DRAWINGS

The various features and advantages of the present invention may be more readily understood with reference to the following detailed description taken in conjunction with the accompanying drawings in which:

FIGURE 1 is an illustration of a right ventricular (RV) pacing electrode and an active pulse generator case placed subcutaneously (SubQ) in the patient's left pectoral region used to sense the far-field ventricular electrogram;

FIGURES 2A and 2B are a flow diagram illustrating the processing of electrograms in order to deliver defibrillation energy when the heart is most susceptible to being defibrillated;

FIGURE 3 is a simulated recording of a far-field electrogram recording during ventricular fibrillation and showing delivery of a defibrillation shock; and

FIGURES 4-9 illustrate various sensing configurations for use with the invention.

DETAILED DESCRIPTION OF THE INVENTION

The present invention is directed toward the provision of a defibrillator that delivers a defibrillation shock to the heart during a period of ventricular fibrillation (VF) in which the shock is likely to be successful in defibrillating the heart. Variation in the state of excitability of the cells in the heart results in a cyclic period of increased susceptibility to defibrillation. The susceptible period occurs when the number of excitable cells is low, i.e., a higher state of total depolarization. The absolute VF voltage (AVFV) is a crude marker of the number of excitable cells.

FIGURE 1 is an illustration of a right ventricular (RV) pacing electrode 10 and an active pulse generator case 12 located subcutaneously (SubQ) in the patient's left pectoral region for sensing the far-field ventricular electrogram. Unlike the bipolar electrogram that would be obtained by sensing from the RV pacing electrode 10 to the RV sensing and/or defibrillation electrode 14 (the usual combination for sensing), the far-field, or unipolar, electrogram provides the necessary information to determine the period of increased susceptibility to defibrillation. A conventional bipolar sensor may also be used to provide ECG signals to the fibrillation detection circuitry.

FIGURES 2A and 2B are a flow diagram of the steps performed by the logic and detection circuitry of the implantable defibrillator for processing the near-field and far-field ventricular electrograms in order to deliver defibrillation energy when the heart is most susceptible to being defibrillated. A fibrillation of the patient's heart is detected at step 30 using conventional detection techniques and a near-field electrogram. Charging of the high voltage (HV) capacitors is started at step 32 following VF detection. A running sequence of windows is started at step 34 to determine the AVFV for the far-field electrogram during capacitor charging. In the preferred embodiment, the running sequence of



windows is about 5 seconds long but it could be longer or shorter. A series of smaller windows, each window in the sequence is about 250 msec long. Only the most recent windows comprising the sequence are used for calculations in setting the threshold to be discussed below. It should be noted that the sequence of windows, which is chosen to have a time long enough to optimally characterize the signal strength or identify a local relative maximum signal, does not have a physical existence but is a continuing calculation of a most recent fixed period of the electrogram signal. In the preferred embodiment, a new window is calculated every 250 msec, thus providing a sequence of slices along the data signal. Alternatively, a new window could be calculated with each new data point so that the window slides along the data signal. With either method, it is preferred that only the most recent portion of the VF data signal be used to calculate a running average or a relative maximum of the AVFV. The AVFV during each window can be calculated in a number of ways including taking an average of the absolute value of all detected points, taking an average of the absolute value of all peaks, or making an area under the waveform curve calculation or using other comparable algorithms. To facilitate this calculation, it is preferred that the detected ECG be digitized early in the processing as is shown in U.S. Patent No. 5,027,814 to Carroll et al., which patent is incorporated herein by reference.

At this point, a loop is started which lasts during the charging of the HV capacitors. An AVFV is calculated for each new window at step 36 using the far-field EGM. A new average or relative maximum AVFV is then calculated at step 38 with each new window in the sequence. For the calculation of the relative maximum, each time a new value is generated, all the values in the running window are compared and the largest becomes the new relative maximum. The presence of ventricular fibrillation is reconfirmed as step 44. If VF is not still present, the shock is aborted at step 46 and charging of the capacitors is halted. If the VF is still present and the capacitors are not fully charged; step 48 causes the system to loop back to step 38 to calculate the next AVFV of the running sequence of windows.

Referring now to FIGURE 2B, once the HV capacitors are fully charged, as a safety measure, a count down timer is started at step 50. Next, an AVFV threshold is set in step 52 based on the average or relative maximum AVFV at that point in time. If an AVFV which exceeds this threshold is not reached by the time the timer times out at step 54, then the presence of a VF is reconfirmed at step 56. If VF is still present, the defibrillation shock is delivered at step 58 synchronous with the

next detected cardiac complex of the electrogram. If VF is not reconfirmed, then the shock is aborted at step 60. In the preferred embodiment, the timer is set for between about 2 and 6 seconds but a shorter or longer timer could be used. In the preferred embodiment, a running average of the AVFV is used at step 52 to set a threshold for delivery of the defibrillation shock. The average continues to be calculated even after capacitor charging is completed. In an alternative embodiment, an AVFV relative maximum is used to set the threshold at step 52. The value of the threshold must be set so that it is not so high that it is unlikely that an AVFV measured subsequent to completion of HV capacitor charging will be as large as the threshold AVFV but it must be sufficiently large that the benefit of the invention is realized. When the threshold is set based on a running average AVFV, a typical multiplier is between about 1.1 and 1.4. When the threshold is set based on a relative maximum AVFV, a typical multiplier for setting the threshold is between about 0.5 and 0.9.

Once the count down timer is started at step 50 a loop is started to find the appropriate time for delivery of the defibrillation shock. Since the running window continues across the time when the capacitors are fully charged, the defibrillation shock may be delivered at the first peak or trough following the charge completion if the AVFV threshold condition is met at step 62. Since the invention works best if the defibrillation shock is delivered synchronous with a relatively large peak, a second threshold is used to determine whether or not to deliver the defibrillation shock into a particular complex at step 64. This second threshold may be a function of the AVFV average or relative maximum value or may be independently set. If a sufficiently large peak is detected at step 64, then VF is reconfirmed at step 56 and the defibrillation shock is delivered at step 58. Depending on the duration of the running sequence of windows, the first threshold may be met and yet the most current cardiac complex be too small to satisfy the second threshold. This prevents delivering the defibrillation shock at a time when the number of excitable cells is greater than desired to benefit from the method of the invention. In that case, delivery of the shock is delayed for a period of a 250 msec window at step 66. Thus, the system will look at several peaks in the new window to find one which exceeds the second threshold. If one is not found within 250 msec, the AVFV from the latest window is used to calculate an updated average or relative maximum AVFV and set a new threshold at step 52.

FIGURE 3 is a simulated recording of a far-field electrogram showing a ventricular fibrillation. Initially, the heart is in a sinus condition and then

fibrillation occurs. Once the fibrillation is detected, which generally takes some minimum number of short cardiac intervals, charging of the high voltage capacitor of the defibrillator is initiated. At that time, the running sequence of windows is initiated. Depending on the fibrillation interval, each window will capture about 1 to 4 cardiac complexes. A longer window of up to one second or more could be used. The digitized fibrillation voltage signal is then processed to provide the AVFV for that window. The figure shows a defibrillation shock being delivered and the signal returning to a sinus condition.

In addition to the sensing electrode configuration shown in FIGURE 1, which uses an RV pacing electrode to an active pulse generator case, far field sensing may be accomplished using electrograms from other combinations of electrodes, including those of FIGURES 4-9. Some electrode configurations may be better than others for determining the relative amount of depolarization of the heart, and the preferred configuration may be different for different patients. To determine the best configuration for a given patient, various combinations of the leads in a patient may be tested at implant to see which is most effective. Alternatively, the sensing configuration of the implantable defibrillator could be externally programmable after implantation. The most effective configuration could be determined during post-operative testing.

FIGURE 4 is a schematic illustration of sensing the far-field ventricular electrogram using an RV pacing electrode 10 and a SubQ defibrillation electrode 16. This is the same configuration shown in FIGURE 1, except that a separate SubQ defibrillation electrode 16 is used instead of an active pulse generator case 12. FIGURE 5 is a schematic illustration of sensing the far-field ventricular electrogram using an RV pacing electrode 10 and an SVC defibrillation electrode 18. FIGURE 6 is a schematic illustration of sensing the far-field ventricular electrogram using an RV pacing electrode 10, a superior vena cava (SVC) defibrillation electrode 18 and a SubQ defibrillation electrode 16. In this configuration, the RV pacing electrode acts as one pole, and the SVC and SubQ electrodes are taken in combination to form the second pole, so that only one far-field ventricular electrogram is produced.

In an alternative embodiment, the RV and SVC electrodes may form one sensing pair, and the RV and SubQ electrodes may form a second sensing pair, thus producing two far-field ventricular electrograms. In this way, additional information is obtained regarding the direction of the fibrillation wavefronts, and vectorcardiography may be used to further determine the period during which a defibrillation shock will be most likely to successfully terminate fibrillation.

Two examples of utilizing two VF signals are shown in FIGURES 6 and 9. As shown in FIGURES 6 and 9, the sensing paths formed by the electrode pairs are approximately perpendicular to each other. In this way, when the fluctuation is great in both signals, the sum of their peak to peak envelope voltages will be greatest, representing the greatest level of organization in the heart. This in turn corresponds to the window of VF susceptible to defibrillation. To add an additional level of control to the timing of shock delivery, during this window just described, the exact moment of delivery may be timed with a relative maximum voltage from one of the signals or from the sum of the signals or from the sum of the rectified signals. Again, the optimum choice may be patient-dependent.

As another example of using the two fibrillation signals, instead of summing the envelope voltages, they can be compared to each other at each point in time, with only the larger peak to peak voltage stored. In this way, two medium peak to peak voltages would not have the same magnitude as one large and one small; the set with the one large signal would be considered to have a higher level of organization, and therefore be more susceptible to defibrillation. As before, the exact moment of energy delivery may be timed to coincide with a absolute VF voltage, this time using the signal with the large envelope voltage.

FIGURE 7 is a schematic illustration of sensing the far-field ventricular electrogram using an RV defibrillation electrode 14 and a SubQ defibrillation electrode 16. FIGURE 8 illustrates sensing the far-field ventricular electrogram using an RV defibrillation electrode 14 and an SVC defibrillation electrode 18.

FIGURE 9 is a schematic illustration of sensing the far-field ventricular electrogram using an RV defibrillation electrode 14, an SVC defibrillation electrode 18 and a SubQ defibrillation electrode 16. As described with reference to FIGURE 6, the RV electrode may be used with the SVC and SubQ electrodes either in combination or separately to form one combined or two separate ventricular electrograms. It is noted that an active pulse generator case may be used for the SubQ electrode in each of the embodiments shown in FIGURES 4, 6, 7 and 9.

It should be understood that various alternatives to the embodiments of the invention described herein may be employed in practicing the invention. It is thus intended that the following claims define the scope of the invention and that structures and methods within the scope of these claims and their equivalents be covered thereby.

Claims

1. An implantable defibrillator, comprising:
 - a pulse generator (12) including a battery, high voltage capacitor means and a charging circuit for charging said capacitor means from said battery;
 - bipolar fibrillation sensing means;
 - unipolar sensing means for monitoring a fibrillation voltage; and
 - electrode means (14) for delivering a defibrillation shock from said capacitor during a period of ventricular fibrillation after said capacitor is charged,
 characterized in that said unipolar sensing means is adapted for determining a relative measure of the amplitude of peaks and troughs of said fibrillation voltage over a period of time following fibrillation detection, and that said defibrillation shock is delivered when a measure of the amplitude of the peaks and troughs of said fibrillation voltage exceeds said relative measure by a desired amount.
2. The defibrillator of claim 1, characterized by a safety timer, a shock being delivered upon timing out of said safety timer if a period of ventricular fibrillation when said measure of the amplitude of the peak and trough voltages exceeds said relative measure by a desired amount has not been reached.
3. The defibrillator of claim 1 or 2, characterized by means for delivering said defibrillation shock synchronous with a peak or trough of said fibrillation voltage having an amplitude exceeding a threshold set as a function of said average amplitude of said peaks and troughs.
4. The defibrillator of any of claims 1 - 3, characterized in that said bipolar fibrillation sensing means comprises a first electrode and a second electrode, and wherein said unipolar sensing means comprises said first electrode and a third electrode different from said second electrode.
5. The defibrillator of claim 4, characterized in that said first electrode comprises either a right ventricular pacing electrode (10) or a right ventricular defibrillation electrode and said third electrode comprises one of a subcutaneous defibrillation electrode (16), an electrically active portion of the implantable defibrillator case, or a superior vena cava defibrillation electrode (18).
6. The defibrillator of any of claims 1 - 5, characterized in that said unipolar sensing means comprises at least three electrodes.
7. The defibrillator of any of claims 4 - 6, characterized in that said electrode means (14) for delivering a defibrillation shock includes said second and third electrodes.
8. An implantable defibrillator, comprising:
 - means for detecting fibrillation in a patient's heart;
 - means for charging a high voltage capacitor in said implantable defibrillator;
 - means for monitoring a fibrillation voltage across first and second implanted electrodes spaced apart from each other by at least 4 centimeters, a waveform of said fibrillation voltage including peaks and troughs;
 - means for delivering a defibrillation shock to said patient's heart following charging of said capacitor during a period of ventricular fibrillation,
 characterized by means for determining relative magnitude of the amplitude of the peaks and troughs of said fibrillation voltage and setting a threshold based on said relative magnitude, said defibrillation shock being delivered when a current value of said relative magnitude of the amplitudes of the peaks and troughs of said fibrillation voltage exceeds said threshold.
9. The defibrillator of claim 8, characterized in that said means for determining said relative magnitude of the amplitude of the peaks and troughs of said fibrillation voltage includes means for calculating an average absolute value amplitude of said peaks and troughs for a running sequence of windows.
10. The defibrillator of claim 8 or 9, characterized in that said means for delivering a defibrillation shock includes means for synchronizing said shock with a peak or trough of said fibrillation voltage having an amplitude exceeding a second threshold set as a function of measured absolute values of said peaks and troughs of said fibrillation voltage from said means for determining a relative magnitude.

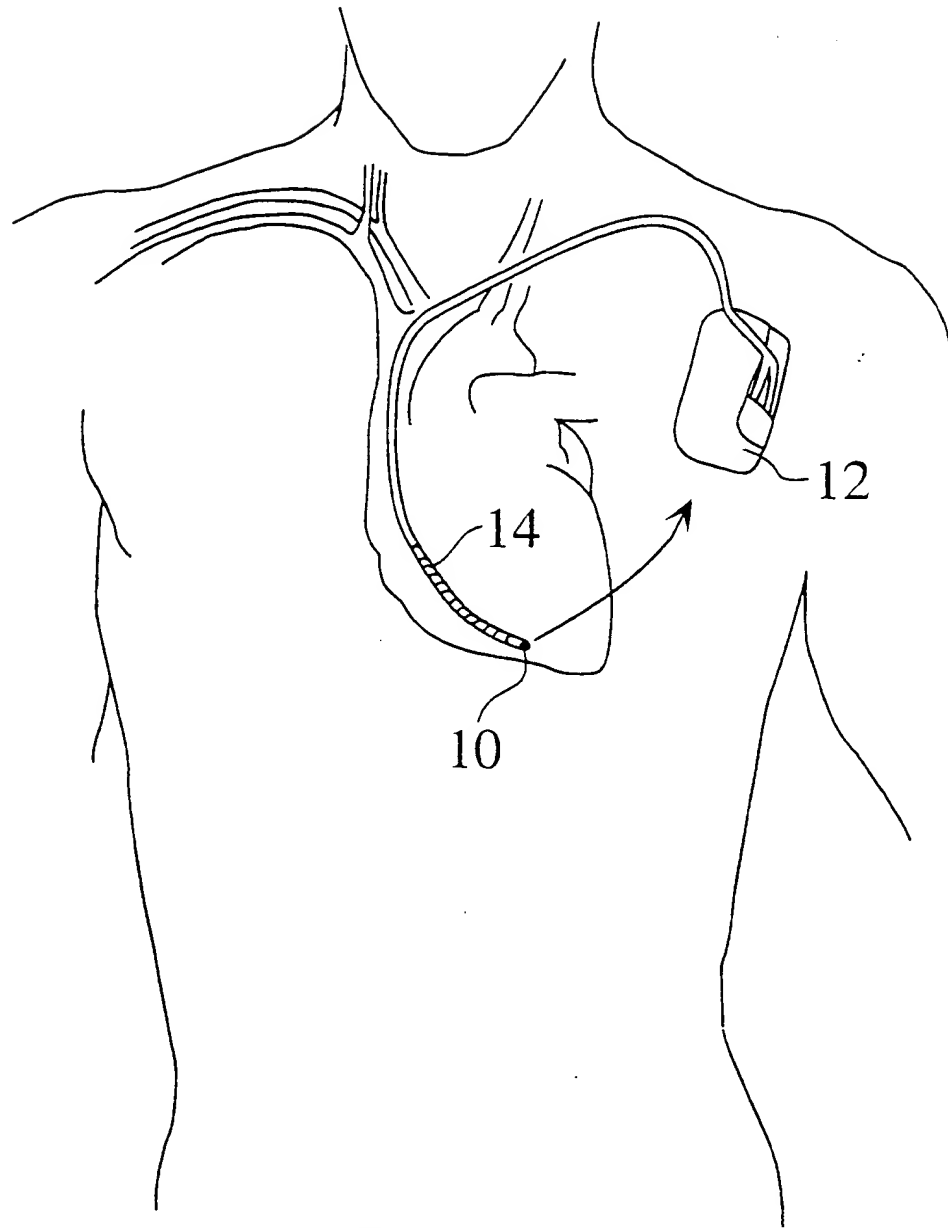


Figure 1

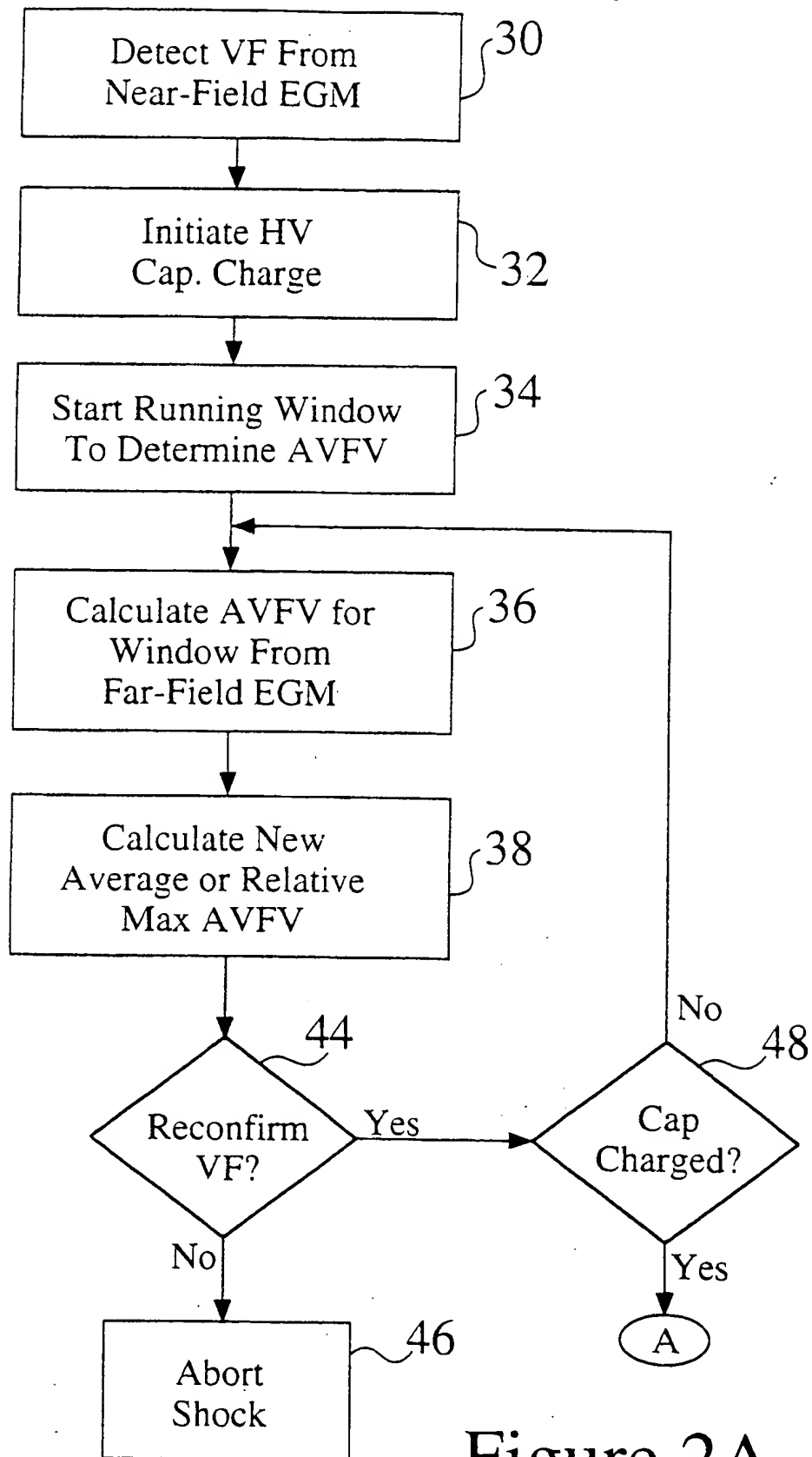


Figure 2A

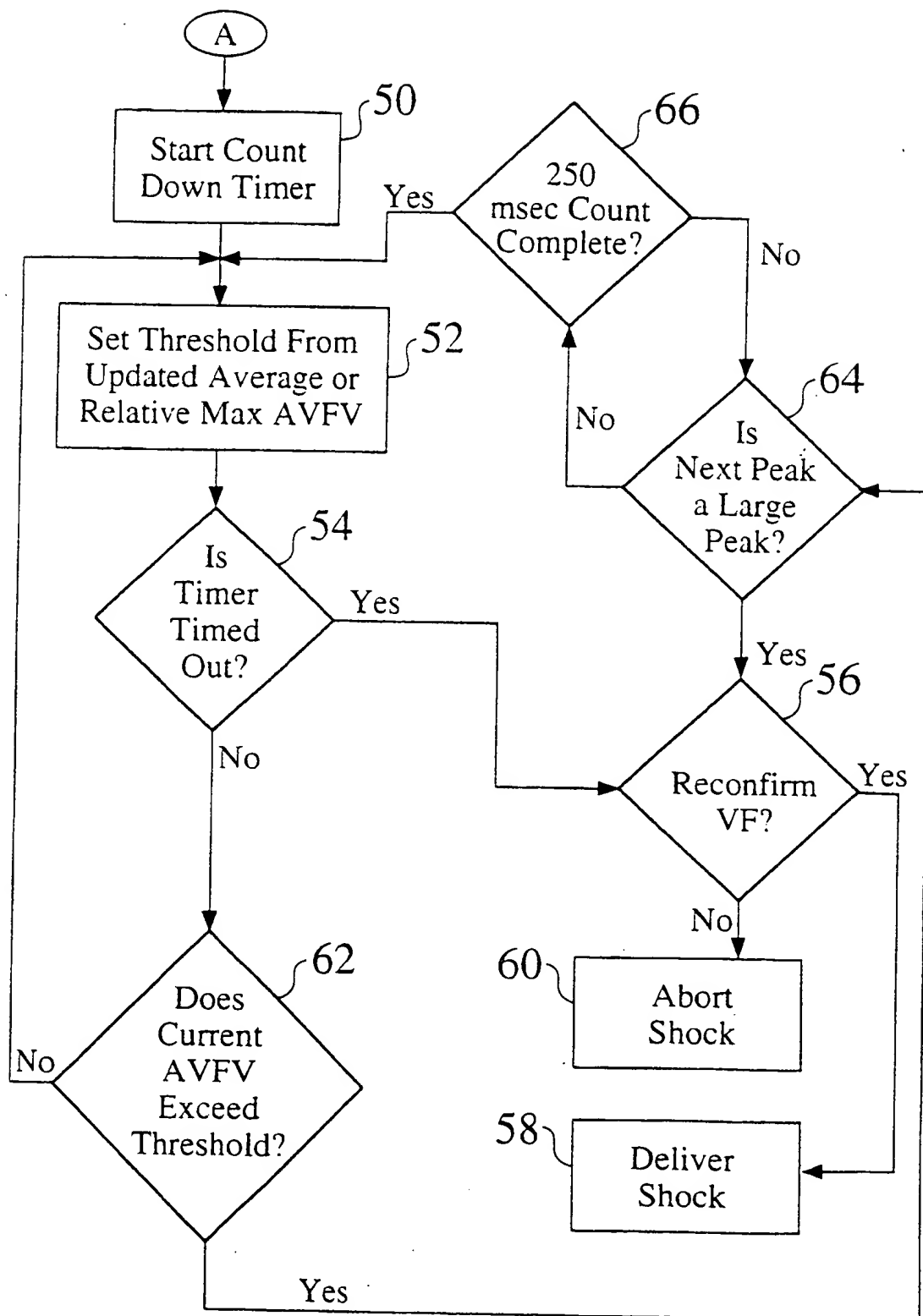


Figure 2B

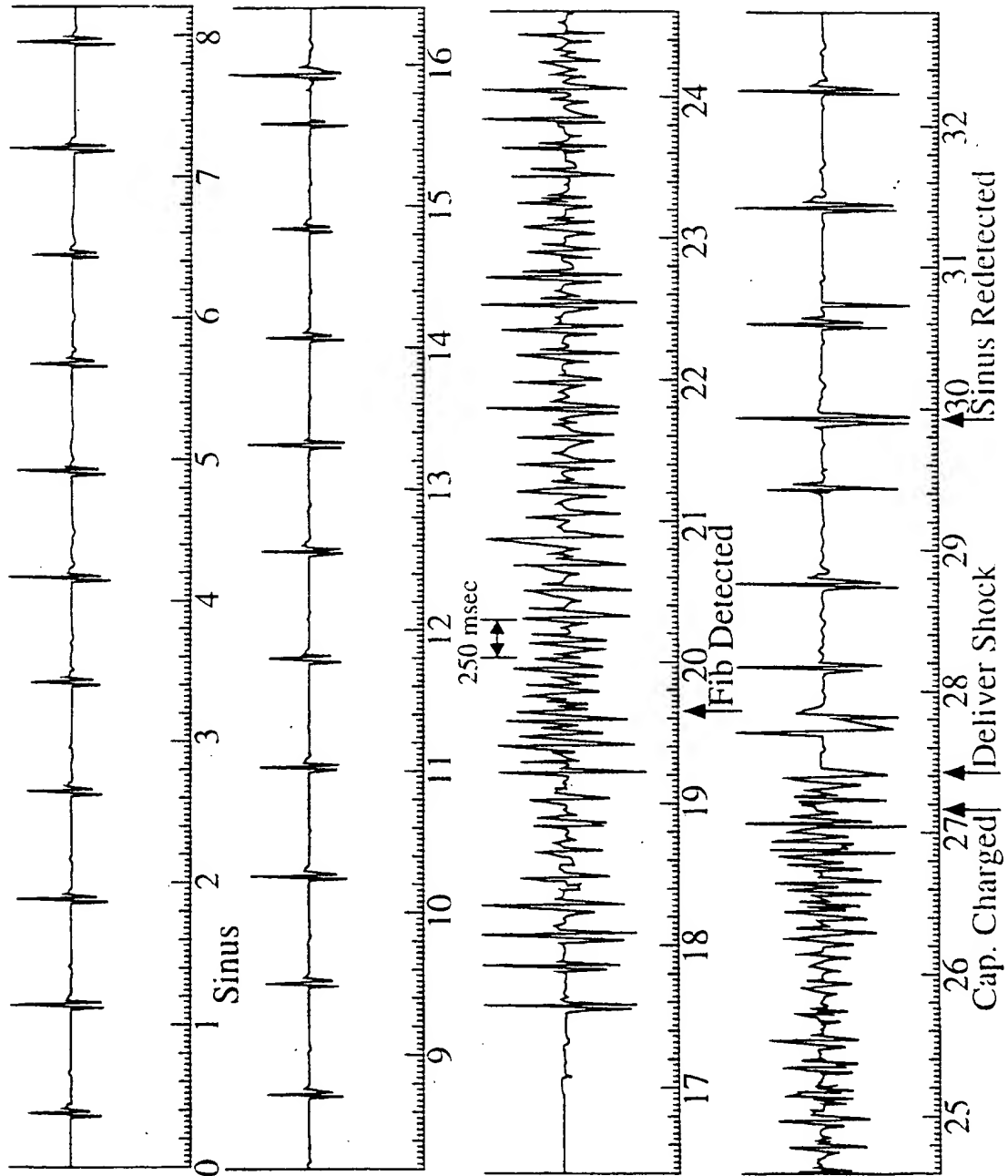


Figure 3

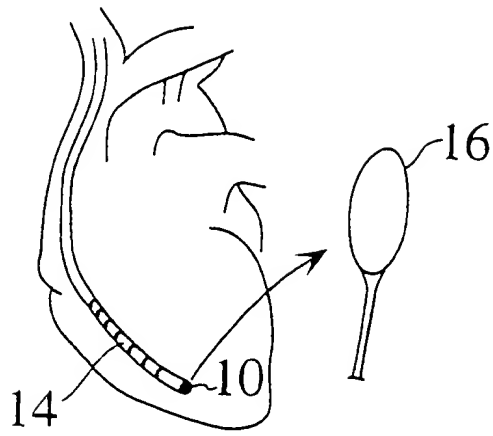


Figure 4

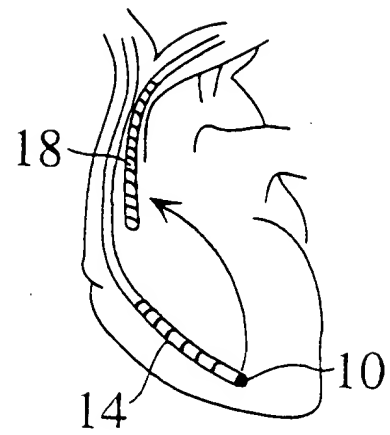


Figure 5

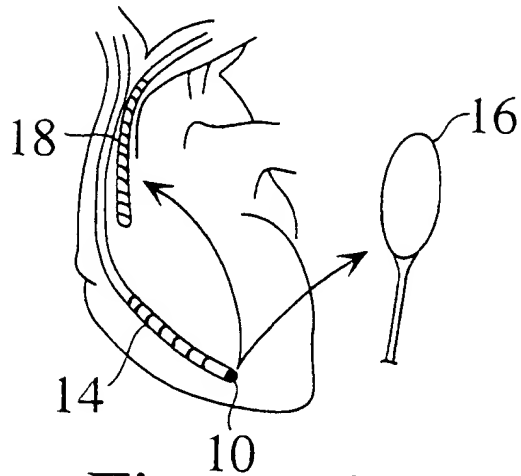


Figure 6

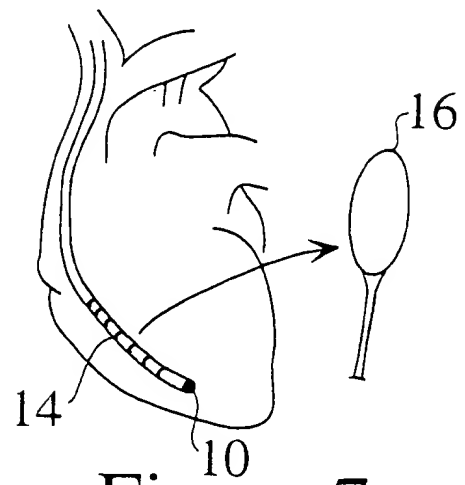


Figure 7

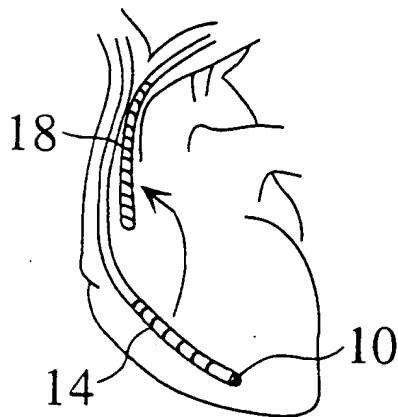


Figure 8

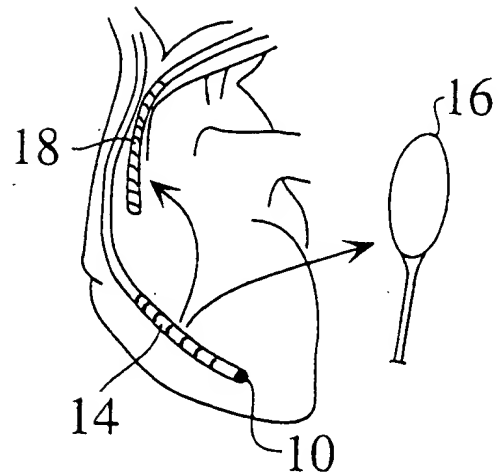


Figure 9

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(71) Applicant: VENTRITEX, INC.
Sunnyvale, California 94025 (US)

(72) Inventor: Fain, Eric S.
Menlo Park, CA 94025 (US)

(74) Representative:
Witte, Alexander, Dr.-Ing. et al
Witte, Weller, Gahlert, Otten & Steil,
Patentanwälte,
Rotebühlstrasse 121
70178 Stuttgart (DE)

(54) Method and apparatus for delivering defibrillation shocks with improved effectiveness

(57) A method for defibrillating a patient's heart with an implantable defibrillator having defibrillation synchronized to the fibrillation waveform is disclosed. The defibrillator is used to detect fibrillation in a patient's heart, monitor at least one far-field fibrillation voltage across two spaced-apart electrodes while the high voltage capacitors are charging, then deliver a high energy shock during a period of ventricular fibrillation when the absolute values of the peak and trough voltages exceed a threshold based on a desired percentage greater than

a running average of the absolute values of the peaks and troughs of the fibrillation voltages monitored during and after capacitor charging. The shock is then delivered synchronous with the next significant cardiac complex. The defibrillator also includes a safety timer so that if the threshold is not reached within a specified safe time limit, the energy shock will be delivered without further delay.

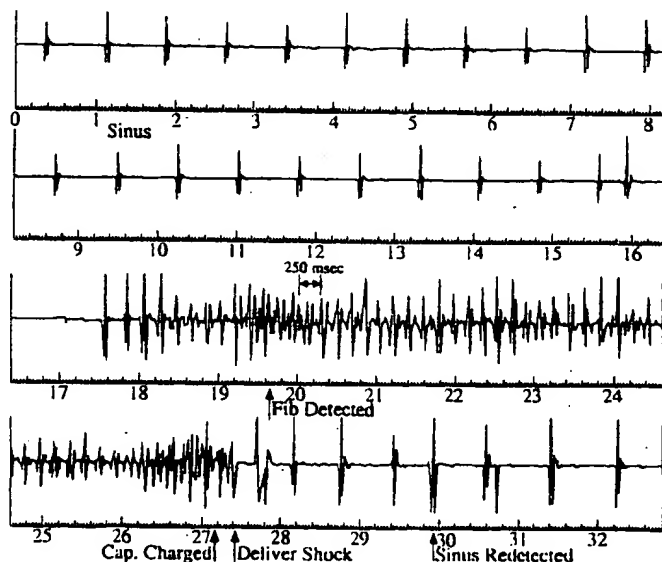


Figure 3

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EUROPEAN SEARCH REPORT

Application Number
EP 95 10 4607

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.6)
A	EP 0 350 160 A (INTERMEDICS) * column 6, line 35 - column 14, line 59 * ---	1,2,4,8	A61N1/39
A	US 4 523 595 A (ZIBELL) * column 3, line 9 - column 6, line 59 * ---	1,2,8,9	
A	FR 2 528 708 A (MIROWSKI) * page 6, line 8 - page 7, line 21 * * page 11, line 15 - page 12, line 26 * ---	1,3,8,10	
A	US 3 612 041 A (RAGSDALE) * the whole document * ---	1,4-6,8,9	
A	WO 81 02665 A (DATAMEDIX) * page 10, line 11 - page 19, line 24 * -----	1,8	
			TECHNICAL FIELDS SEARCHED (Int.Cl.6)
			A61N
The present search report has been drawn up for all claims			
Place of search		Date of completion of the search	Examiner
THE HAGUE		9 December 1997	Lemerrier, D
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